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(FILE 'HOME' ENTERED AT 11:43:22 ON 21 MAR 2008)

FILE 'CAPLUS' ENTERED AT 11:43:32 ON 21 MAR 2008

E US2005-550760/APPS

L1 1 S E3  
SEL L1 RN

FILE 'REGISTRY' ENTERED AT 11:44:52 ON 21 MAR 2008

L2 39 S E1-E39

FILE 'STNGUIDE' ENTERED AT 11:46:11 ON 21 MAR 2008

FILE 'REGISTRY' ENTERED AT 12:00:33 ON 21 MAR 2008

E "1H-IMIDAZOLE-5-CARBOXYLIC ACID, 2-BUTYL-4-CHLORO-1-((2'-(2H-

E "1H-IMIDAZOLE-5-CARBOXYLIC ACID, 2-BUTYL-4-(PENTAFLUOROETHYL)

E "3H-IMIDAZO(4,5-B)PYRIDINE, 2-ETHYL-5,7-DIMETHYL-3-((2'-(2H-T

E "5-PYRIMIDINECARBOXYLIC ACID, 4-(BUTYL((2'-(2H-TETRAZOL-5-YL)

E "1H-TETRAZOLE, 5-(4'-((3,5-DIBUTYL-1H-1,2,4-TRIAZOL-1-YL)METH

E "2H-IMIDAZOL-2-ONE, 1,4-DIBUTYL-1,3-DIHYDRO-3-((2'-(1H-TETRAZ

E "QUINOLINE, 2-ETHYL-4-((2'-(2H-TETRAZOL-5-YL)(1,1'-BIPHENYL)-

E "QUINOLINE, 2-ETHYL-5,6,7,8-TETRAHYDRO-4-((2'-(2H-TETRAZOL-5-

L3 1 S "1H-IMIDAZOLE-5-CARBOXYLIC ACID, 2-BUTYL-4-CHLORO-1-((2'-(2H-

L4 1 S "1H-IMIDAZOLE-5-CARBOXYLIC ACID, 2-BUTYL-4-(PENTAFLUOROETHYL)

L5 1 S "3H-IMIDAZO(4,5-B)PYRIDINE, 2-ETHYL-5,7-DIMETHYL-3-((2'-(2H-T

L6 1 S "5-PYRIMIDINECARBOXYLIC ACID, 4-(BUTYL((2'-(2H-TETRAZOL-5-YL)

L7 1 S "1H-TETRAZOLE, 5-(4'-((3,5-DIBUTYL-1H-1,2,4-TRIAZOL-1-YL)METH

L8 1 S "2H-IMIDAZOL-2-ONE, 1,4-DIBUTYL-1,3-DIHYDRO-3-((2'-(1H-TETRAZ

L9 1 S "QUINOLINE, 2-ETHYL-4-((2'-(2H-TETRAZOL-5-YL)(1,1'-BIPHENYL)-

L10 1 S "QUINOLINE, 2-ETHYL-5,6,7,8-TETRAHYDRO-4-((2'-(2H-TETRAZOL-5-

FILE 'CAPLUS, USPATFULL, USPATOLD, USPAT2' ENTERED AT 12:05:44 ON 21 MAR  
2008

L11 520 S L3-L10

L12 38368 S (METABOLIC SYNDROME X) OR (SYNDROME X) OR (INSULIN RESISTANCE

L13 12 S L11 AND L12

L14 4 S L13 AND PY<2004

SAVE TEMP ALL A10550760/L

=> d ibib abs hitstr 1-4

L14 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:701277 CAPLUS <<LOGINID::20080321>>

DOCUMENT NUMBER: 140:104847

TITLE: Effects of angiotensin II receptor antagonists on  
insulin resistance syndrome and  
leptin in sucrose-fed spontaneously hypertensive rats

AUTHOR(S): Umeda, Mamoru; Kanda, Tsugiyasu; Murakami, Masami

CORPORATE SOURCE: Department of Laboratory Medicine Gunma University  
School of Medicine, Maebashi, Japan

SOURCE: Hypertension Research (2003), 26(6), 485-492

CODEN: HRESE4; ISSN: 0916-9636

PUBLISHER: Japanese Society of Hypertension

DOCUMENT TYPE: Journal

LANGUAGE: English

AB In order to investigate the usefulness of angiotensin II type 1 receptor (AT1) antagonists (ARA) in the treatment of hypertension with insulin resistance syndrome, we studied the effects of a high dose sucrose diet and ARA on insulin sensitivity, plasma lipids, and leptin in spontaneous hypertensive rats (SHR) and WistarKyoto rats (WKY). SHR and WKY were divided into three groups and treated for 12 wk: those fed a standard chow, those given a sucrose-rich chow or those given a sucrose-rich chow and ARA. While in SHR the weight of both s.c. and mesenteric adipose tissue was greater in the sucrose-rich chow fed animals than in the standard chow fed animals, ARA treatment significantly decreased the wts. of both s.c. and mesenteric adipose tissue. ARA treatment decreased free fatty acid and triglyceride in SHR, and increased high d. lipoprotein cholesterol in SHR and WKY. Homeostasis model assessment-insulin resistance (HOMA-IR) index, plasma levels of leptin, and leptin mRNA in mesenteric adipose tissue were significantly greater in the sucrose-rich chow fed animals than in the standard chow fed animals, and significantly lower in the ARA-treated sucrose-rich chow fed animals than in the sucrose-rich chow fed animals in both SHR and WKY. ARA improved insulin resistance, and reduced plasma leptin and leptin mRNA in adipose tissue. These results suggest that the improvement of insulin resistance by ARA may be attributed, at least in part, to the reduction of adipose tissue weight. It is concluded that ARA is useful in the treatment of patients with hypertension and concomitant insulin resistance syndrome.

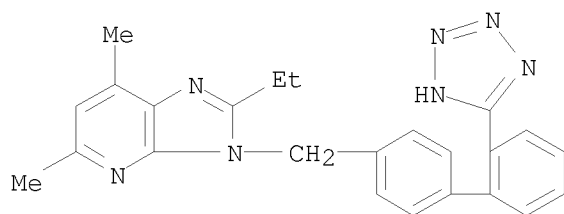
IT 133240-46-7, L-158809

RL: DMA (Drug mechanism of action); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(effects of angiotensin II receptor antagonists on insulin resistance syndrome and leptin in sucrose-fed spontaneously hypertensive rats)

RN 133240-46-7 CAPLUS

CN 3H-Imidazo[4,5-b]pyridine, 2-ethyl-5,7-dimethyl-3-[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (CA INDEX NAME)



REFERENCE COUNT: 43 THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 2 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2002:214313 USPATFULL <<LOGINID::20080321>>

TITLE: Use of an angiotensin II receptor antagonist for the preparation of drugs to increase the survival rate of renal transplant patients

INVENTOR(S): Remuzzi, Giuseppe, Bergamo, ITALY

PATENT ASSIGNEE(S): MERCK SHARP & DOHME (Italia) S.p.A., Roma, ITALY (non-U.S. corporation)

	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 2002115702	A1	20020822	<--
	US 6576652	B2	20030610	
APPLICATION INFO.:	US 2002-76396	A1	20020219	(10)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 2000-509791, filed on 30 Mar 2000, PENDING			

	NUMBER	DATE
PRIORITY INFORMATION:	IT 1997-RM586	19970930
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	BROWDY AND NEIMARK, P.L.L.C., SUTIE 300, 624 NINTH STREET, N.W., WASHINGTON, DC, 20001-5303	
NUMBER OF CLAIMS:	21	
EXEMPLARY CLAIM:	1	
LINE COUNT:	2068	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to the use, for the preparation of drugs to increase the survival rate of transplant patients, including renal and heart transplant patients, of a therapeutically effective amount of an angiotension II receptor antagonist compound, such as the class of substituted imidazoles represented by formula (I) and in particular by losartan potassium, 2-butyl-4-chloro-[(2'-tetrazol-5-yl)biphenyl-4-yl]methyl]-5-(hydroxymethyl)imidazole potassium salt.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

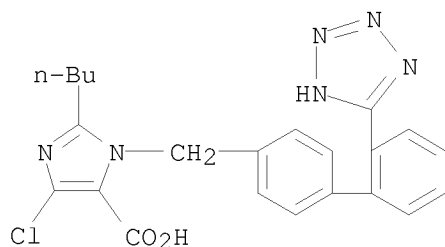
IT 124750-92-1P

(preparation of drugs to increase the survival rate of renal transplant patients)

RN 124750-92-1 USPATFULL

CN 1H-Imidazole-5-carboxylic acid, 2-butyl-4-chloro-1-[[2'-(2H-tetrazol-5-

yl)[1,1'-biphenyl]-4-yl)methyl]- (CA INDEX NAME)



L14 ANSWER 3 OF 4 USPATFULL on STN

ACCESSION NUMBER: 1999:121402 USPATFULL &lt;&lt;LOGINID::20080321&gt;&gt;

TITLE: Insulin sensitivity with angiotensin II receptor blocking imidazoles

INVENTOR(S): Eide, Ivar K., P.O. Box 2000, Rahway, NJ, United States 07065

Moan, Andreas, P.O. Box 2000, Rahway, NJ, United States 07065

Kjeldsen, Sverre E., P.O. Box 2000, Rahway, NJ, United States 07065

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5962500		19991005 <--
APPLICATION INFO.:	US 1998-128138		19980803 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1997-958236, filed on 27 Oct 1997, now abandoned which is a continuation of Ser. No. US 1996-775696, filed on 31 Dec 1996, now abandoned which is a continuation of Ser. No. US 1995-406620, filed on 20 Mar 1995, now abandoned		

	NUMBER	DATE
PRIORITY INFORMATION:	GB 1994-6573	19940331
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Cooney, Jr., John M.	
LEGAL REPRESENTATIVE:	Camara, Valerie J., Daniel, Mark R.	
NUMBER OF CLAIMS:	7	
EXEMPLARY CLAIM:	1	
LINE COUNT:	1510	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention relates to a novel method of using an Angiotensin II antagonist for the improvement of insulin sensitivity alone or in conjunction with the treatment of hypertension. Angiotensin II antagonists such as the class of substituted imidazoles represented by formula I: ##STR1## and specifically by Losartan, 2-butyl-4-chloro-1-[(2'-tetrazol-5-yl)biphenyl-4-yl)methyl]-5-(hydroxymethyl)imidazole potassium salt.

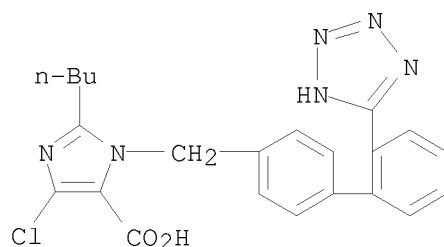
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 124750-92-1P

(improvement of insulin sensitivity with angiotensin II  
receptor-blocking imidazoles)

RN 124750-92-1 USPATFULL

CN 1H-Imidazole-5-carboxylic acid, 2-butyl-4-chloro-1-[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (CA INDEX NAME)



L14 ANSWER 4 OF 4 USPAT2 on STN

ACCESSION NUMBER: 2002:214313 USPAT2 <<LOGINID::20080321>>

TITLE: Use of an angiotensin II receptor antagonist for the  
preparation of drugs to increase the survival rate of  
renal transplant patients

INVENTOR(S): Remuzzi, Giuseppe, Bergamo, ITALY

PATENT ASSIGNEE(S): Merck Sharp & Dohme (Italia) S.p.A., Rome, ITALY  
(non-U.S. corporation)

	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 6576652	B2	20030610	<--
APPLICATION INFO.:	US 2002-76396		20020219	(10)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 2000-509791, filed on 30 Mar 2000, now abandoned			

	NUMBER	DATE
PRIORITY INFORMATION:	IT 1997-RM586	19970930
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Weddington, Kevin E.	
LEGAL REPRESENTATIVE:	Browdy & Neimark PLLC	
NUMBER OF CLAIMS:	21	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	0 Drawing Figure(s); 0 Drawing Page(s)	
LINE COUNT:	2179	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to the use, for the preparation of drugs to increase the survival rate of transplant patients, including renal and heart transplant patients, of a therapeutically effective amount of an angiotensin II receptor antagonist compound, such as the class of substituted imidazoles represented by formula (I) and in particular by losartan potassium, 2-butyl-4-chloro-[(2'-tetrazol-5-yl)biphenyl-4-

il]methyl]-5-(hydroxymethyl)imidazole potassium salt.

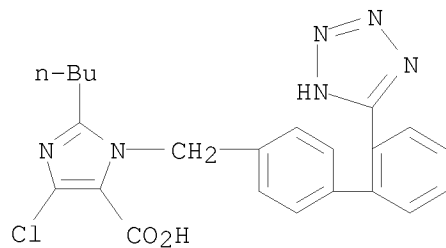
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 124750-92-1P

(preparation of drugs to increase the survival rate of renal transplant patients)

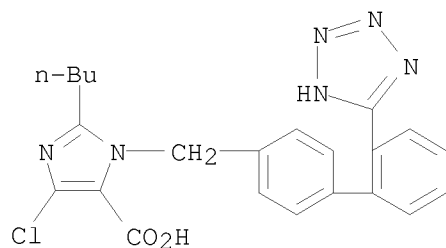
RN 124750-92-1 USPAT2

CN 1H-Imidazole-5-carboxylic acid, 2-butyl-4-chloro-1-[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (CA INDEX NAME)



=> d 13

L3 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
RN 124750-92-1 REGISTRY  
ED Entered STN: 12 Jan 1990  
CN 1H-Imidazole-5-carboxylic acid, 2-butyl-4-chloro-1-[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (CA INDEX NAME)  
OTHER CA INDEX NAMES:  
CN 1H-Imidazole-5-carboxylic acid, 2-butyl-4-chloro-1-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (9CI)  
OTHER NAMES:  
CN E 3174  
CN EXP 3174  
MF C22 H21 Cl N6 O2  
CI COM  
SR CA  
LC STN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN\*, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CHEMCATS, DDFU, DRUGU, EMBASE, IMSRESEARCH, IPA, MEDLINE, RTECS\*, SYNTHLINE, TOXCENTER, USPAT2, USPATFULL  
(\*File contains numerically searchable property data)

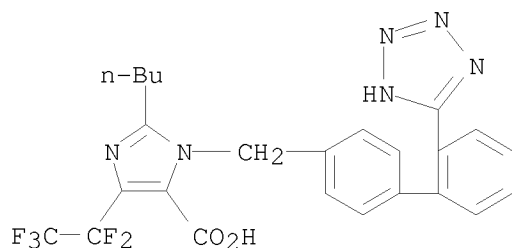


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

260 REFERENCES IN FILE CA (1907 TO DATE)  
7 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
260 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d 14

L4 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
RN 124750-94-3 REGISTRY  
ED Entered STN: 12 Jan 1990  
CN 1H-Imidazole-5-carboxylic acid, 2-butyl-4-(pentafluoroethyl)-1-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (9CI) (CA INDEX NAME)  
MF C24 H21 F5 N6 O2  
SR CA  
LC STN Files: CA, CAPLUS, USPATFULL

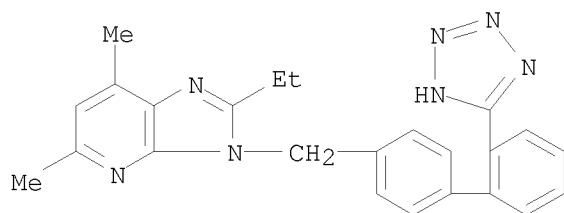


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1907 TO DATE)  
 1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
 5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d 15

L5 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
 RN 133240-46-7 REGISTRY  
 ED Entered STN: 12 Apr 1991  
 CN 3H-Imidazo[4,5-b]pyridine, 2-ethyl-5,7-dimethyl-3-[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (CA INDEX NAME)  
 OTHER CA INDEX NAMES:  
 CN 3H-Imidazo[4,5-b]pyridine, 2-ethyl-5,7-dimethyl-3-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (9CI)  
 OTHER NAMES:  
 CN L 158809  
 MF C24 H23 N7  
 SR CA  
 LC STN Files: ADISINSIGHT, BEILSTEIN\*, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, DDFU, DRUGU, EMBASE, IPA, PHAR, RTECS\*, TOXCENTER, USPAT2, USPATFULL  
 (\*File contains numerically searchable property data)



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

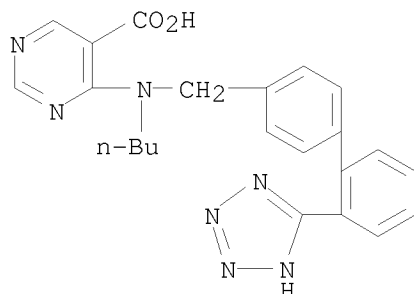
173 REFERENCES IN FILE CA (1907 TO DATE)  
 6 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA



## 173 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=&gt; d 16

L6 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
RN 141872-46-0 REGISTRY  
ED Entered STN: 19 Jun 1992  
CN 5-Pyrimidinecarboxylic acid, 4-[butyl[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]amino]- (CA INDEX NAME)  
OTHER CA INDEX NAMES:  
CN 5-Pyrimidinecarboxylic acid, 4-[butyl[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]amino]- (9CI)  
OTHER NAMES:  
CN Abbott 81282  
MF C23 H23 N7 O2  
CI COM  
SR CA  
LC STN Files: BEILSTEIN\*, BIOSIS, CA, CAPLUS, CASREACT, CHEMCATS, IMSRESEARCH, MEDLINE, PROUDDR, TOXCENTER, USPATFULL  
(\*File contains numerically searchable property data)



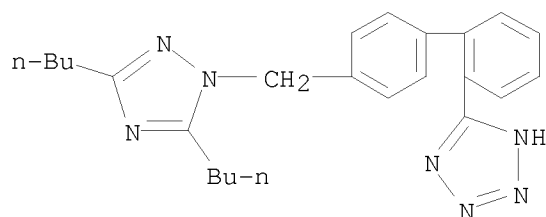
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12 REFERENCES IN FILE CA (1907 TO DATE)  
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12 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=&gt; d 17

L7 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
RN 140120-42-9 REGISTRY  
ED Entered STN: 03 Apr 1992  
CN 1H-Tetrazole, 5-[4'-[(3,5-dibutyl-1H-1,2,4-triazol-1-yl)methyl][1,1'-biphenyl]-2-yl]- (9CI) (CA INDEX NAME)  
OTHER NAMES:  
CN SC 50560  
MF C24 H29 N7  
SR CA

LC STN Files: ADISINSIGHT, CA, CAPLUS, PHAR, PROUSDDR, TOXCENTER, USPATFULL



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

18 REFERENCES IN FILE CA (1907 TO DATE)  
2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
18 REFERENCES IN FILE CAPLUS (1907 TO DATE)

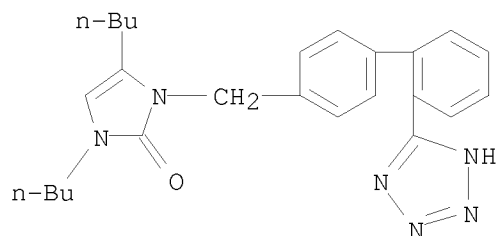
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L8 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
RN 141386-89-2 REGISTRY  
ED Entered STN: 15 May 1992  
CN 2H-Imidazol-2-one, 1,4-dibutyl-1,3-dihydro-3-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN SC 51895  
MF C25 H30 N6 O  
SR CA

LC STN Files: ADISINSIGHT, CA, CAPLUS, PHAR, PROUSDDR, TOXCENTER, USPATFULL

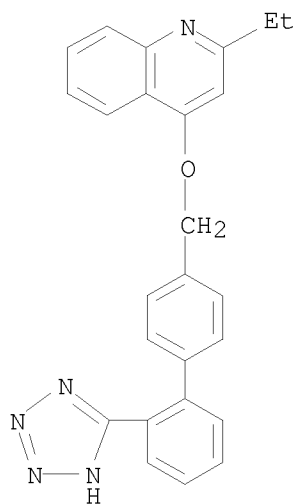


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

12 REFERENCES IN FILE CA (1907 TO DATE)  
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
12 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d 19

L9 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
RN 143494-72-8 REGISTRY  
ED Entered STN: 18 Sep 1992  
CN Quinoline, 2-ethyl-4-[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methoxy]- (CA INDEX NAME)  
OTHER CA INDEX NAMES:  
CN Quinoline, 2-ethyl-4-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methoxy]- (9CI)  
OTHER NAMES:  
CN D 8731  
CN ICI-D 8731  
MF C25 H21 N5 O  
CI COM  
SR CA  
LC STN Files: CA, CAPLUS, CASREACT, MEDLINE, TOXCENTER, USPAT2, USPATFULL



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

35 REFERENCES IN FILE CA (1907 TO DATE)  
2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
35 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d 110

L10 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN  
RN 138620-04-9 REGISTRY  
ED Entered STN: 31 Jan 1992  
CN Quinoline, 2-ethyl-5,6,7,8-tetrahydro-4-[[2'-(2H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methoxy]- (CA INDEX NAME)

## OTHER CA INDEX NAMES:

CN Quinoline, 2-ethyl-5,6,7,8-tetrahydro-4-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methoxy]- (9CI)

## OTHER NAMES:

CN ICI-D 6888

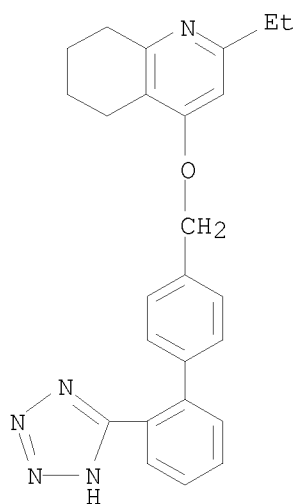
CN ZD 6888

MF C25 H25 N5 O

CI COM

SR CA

LC STN Files: ADISINSIGHT, BIOSIS, CA, CAPLUS, PHAR, PROUSDDR, TOXCENTER, USPAT2, USPATFULL



L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN  
 ACCESSION NUMBER: 2004:857382 CAPLUS <<LOGINID::20080321>>  
 DOCUMENT NUMBER: 141:325747  
 TITLE: Use of an angiotensin II type 1 receptor antagonist  
 for the treatment or prevention of metabolic syndrome  
 INVENTOR(S): Ljunggren, Anders; Svensson, Anders  
 PATENT ASSIGNEE(S): Astrazeneca AB, Swed.  
 SOURCE: PCT Int. Appl., 30 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004087136	A1	20041014	WO 2004-SE505	20040331
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004226517	A1	20041014	AU 2004-226517	20040331
AU 2004226517	B2	20080124		
CA 2520960	A1	20041014	CA 2004-2520960	20040331
EP 1613309	A1	20060111	EP 2004-724927	20040331
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK				
BR 2004008979	A	20060404	BR 2004-8979	20040331
CN 1771033	A	20060510	CN 2004-80009394	20040331
JP 2006522115	T	20060928	JP 2006-507997	20040331
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US 2006194856	A1	20060831	US 2005-550760	20050927 <--
PRIORITY APPLN. INFO.:			SE 2003-988	A 20030403
			WO 2004-SE505	A 20040331
OTHER SOURCE(S):	MARPAT 141:325747			
REFERENCE COUNT:	3	THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT		